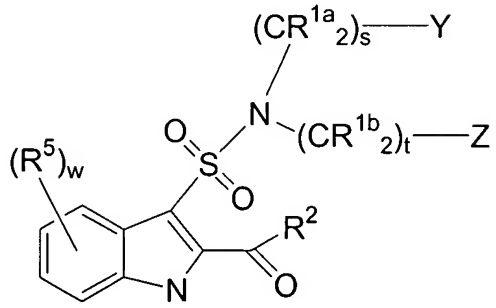
In the claims:

1. (Original) A compound of Formula I:



I

wherein:

R^{1a} and R^{1b} are independently selected from:

1) hydrogen,
2) unsubstituted or substituted C_1-C_{10} alkyl,
3) OR^3,
4) $N(R^3)_2$,
5) unsubstituted or substituted aryl,
6) unsubstituted or substituted heterocycle, and
7) unsubstituted or substituted C_3-C_{10} cycloalkyl;

R^{1c} is independently selected from:

1) hydrogen,
2) C_1-C_{10} alkyl,
3) OR^3,
4) $N(R^3)_2$,
5) C_3-C_{10} cycloalkyl,
6) aryl, and
7) heterocycle;

said alkyl, cycloalkyl, aryl and heterocycle is optionally substituted with at least one substituent selected from R^7;

R^2 is independently selected from:

1) hydrogen,
2) unsubstituted or substituted C_1-C_{10} alkyl,
3) $N(R^3)_2$,
4) OR^3,
5) unsubstituted or substituted aryl, and
6) unsubstituted or substituted C_3-C_{10} cycloalkyl;

R^3 is independently selected from:

1) hydrogen,
2) C_1-C_{10} alkyl,
3) aryl,
4) heterocycle,
5) C_3-C_{10} cycloalkyl,
6) CF_3,
7) C_2-C_6 alkenyl,
8) C_2-C_6 alkynyl,
9) $S(O)_mR^6$, and
10) $C(O)R^6$;

said alkyl, cycloalkyl, aryl, heterocycle, alkynyl, and alkenyl is optionally substituted with at least one substituent selected from R^7;

R^5 is independently selected from:

1) hydrogen,
2) halogen,
3) $-(CR^{1c}{}_2)_nOR^3$,
4) $-(CR^{1c}{}_2)_nR^6$,
5) $-C(O)OR^3$,
6) $-C(O)R^3$,
7) $-C\equiv CR^3$,

8) $-R^3C=C(R^3)_2$,
9) $-OS(O)_mR^6$,
10) $-NO_2$,
11) $-(CR^{1c}{}_2)_nN(R^3)_2$,
12) $-N(R^3)C(O)R^3$,
13) $-N(R^3)S(O)_mR^6$,
14) $-(CR^{1c}{}_2)_nNR^3(CR^{1c}{}_2)_nC(O)NR^3{}_2$,
15) $-O(CR^{1c}{}_2)_nC(O)N(R^3)_2$,
16) $-O(CR^{1c}{}_2)_nC(O)OR^3$,
17) $-NR^3(CR^{1c}{}_2)_nN(R^3)_2$,
18) $-(CR^{1c}{}_2)_nNR^3R^6OR^3$,
19) $-S(O)_mR^6$,
20) $-S(O)_mN(R^3)_2$,
21) -CN,
22) $-(CR^{1c}{}_2)_nN(R^3)(CR^{1c}{}_2)_nR^6$, and
23) $-(CR^{1c}{}_2)_nC(O)N(R^3)_2$;

R^6 is independently selected from:

1) C_1-C_{10} alkyl,
2) C_3-C_{10} cycloalkyl,
3) aryl, and
4) heterocycle;

said, alkyl, cycloalkyl, aryl and heterocycle is optionally substituted with at least one substituent selected from R^7;

R^7 is independently selected from:

1) hydrogen,
2) unsubstituted or substituted C_1-C_{10} alkyl,
3) unsubstituted or substituted C_3-C_{10} cycloalkyl,
4) unsubstituted or substituted aryl,
5) halogen,
6) OR^3,

7) CF_3,
8) unsubstituted or substituted heterocycle,
9) $S(O)_mN(R^3)_2$,
10) $C(O)OR^3$,
11) $C(O)R^3$,
12) CN,
13) $C(O)N(R^3)_2$,
14) $N(R^3)C(O)R^3$,
15) $S(O)_mR^6$, and
16) NO_2;

Y and Z are independently selected from:

1) hydrogen,
2) R^6,
3) OR^3,
4) $N(R^3)_2$,
5) $C(O)OR^3$,
6) $C(O)N(R^3)_2$,
7) $C(O)R^3$,
8) halogen,
9) $N(R^3)(CR^{1c}{}_2)_nC(O)N(R^3)_2$,
10) $S(O)_mN(R^3)_2$,
11) $N(R^3)C(O)OR^3$,
12) $N(R^3)S(O)_mR^6$,
13) $N(R^3)C(O)R^3$,
14) $N(R^3)(CR^{1c}{}_2)_nR^3$,
15) $S(O)_mR^6$,
16) $R^6S(O)_mN(R^3)_2$,
17) $R^6S(O)_mR^6$,
18) $N(R^3)\ S(O)_m\ (CR^{1c}{}_2)_nR^6$,
19) $N(R^3)S(O)_mR^6OR^3$,
20) $N(R^3)C(O)N(R^3)_2$,

21) $N(R^3)C(O)R^6OR^3$,
22) $N(R^3)(CR^{1c}{}_2)_nR^6OR^3$,
23) $N(R^3)OR^3$, and
24) $N(R^3)S(O)_mR^6NO_2$;

m is independently 0, 1 or 2;
n is independently 0 to 6;
s is 0 to 6;
t is 0 to 6;
w is 0 to 4;

or a pharmaceutically acceptable salt or stereoisomer thereof.

2. (Original) The compound according to Claim 1,

wherein:

R^{1a} and R^{1b} are independently selected from:

1) hydrogen,
2) unsubstituted or substituted C_1-C_{10} alkyl,
3) unsubstituted or substituted aryl,
4) unsubstituted or substituted heterocycle, and
5) OR^3;

R^{1c} is independently selected from:

1) hydrogen,
2) C_1-C_{10} alkyl,
3) OR^3,
4) $N(R^3)_2$,
5) aryl, and
6) heterocycle;

said alkyl, aryl and heterocycle is optionally substituted with at least one substituent selected from R^7;

R^2 is:

1) H,

2) unsubstituted or substituted alkyl,
3) OR^3, or
4) $N(R^3)_2$;

R^3 is independently selected from:

1) hydrogen,
2) C_1-C_{10} alkyl,
3) aryl,
4) heterocycle,
5) C_3-C_{10} cycloalkyl,
6) CF_3,
7) $S(O)_mR^6$, and
8) $C(O)R^6$;

said alkyl, cycloalkyl, aryl and heterocycle is optionally substituted with at least one substituent selected from R^7;

R^5 is independently selected from:

1) hydrogen,
2) halogen,
3) $-OR^3$,
4) $-C(O)OR^3$,
5) $-C(O)R^3$,
6) $-C\equiv CR^3$,
7) $-R^3C=C(R^3)_2$,
8) $-OS(O)_mR^6$,
9) $-NO_2$,
10) $-N(R^3)_2$,
11) $-N(R^3)C(O)R^3$,
12) $-N(R^3)S(O)_mR^6$,
13) $-(CR^{1c}{}_2)_nNR^3(CR^{1c}{}_2)_nC(O)NR^3{}_2$,
14) $-O(CR^{1c}{}_2)_nC(O)N(R^3)_2$,
15) $-O(CR^{1c}{}_2)_nC(O)OR^3$,

16) $-NR^3(CR^{1c}_2)_nN(R^3)_2$,
17) $-(CR^{1c}_2)_nNR^3R^6OR^3$,
18) $-S(O)_mR^6$,
19) $-S(O)_mN(R^3)_2$,
20) -CN, and
21) $-(CR^{1c}_2)_nN(R^3)(CR^{1c}_2)_nR^6$;

or a pharmaceutically acceptable salt or stereoisomer thereof.

3. (Original) The compound according to Claim 2,

wherein:

R^{1a} and R^{1b} are independently selected from hydrogen, unsubstituted or substituted C_1-C_{10} alkyl, OR^3, and unsubstituted or substituted aryl;

R^{1c} is independently selected from:

1) hydrogen,
2) C_1-C_{10} alkyl,
3) OR^3, and
4) aryl;

said alkyl and aryl is optionally substituted with at least one substituent selected from R^7;

R^2 is:

1) OR^3, or
2) $N(R^3)_2$;

R^5 is independently selected from:

1) hydrogen,
2) $(CR^{1c}_2)_nR^6$,
3) halogen,
4) $-(CR^{1c}_2)_nOR^3$,
5) $-C(O)OR^3$,
6) $-C(O)R^3$,

7) $-C\equiv CR^3$,
8) $-R^3C=C(R^3)_2$,
9) $(CR^{1c}{}_2)_nC(O)N(R^3)_2$, and
10) $(CR^{1c}{}_2)_nN(R^3)_2$;

Y is:

1) hydrogen,
2) R^6,
3) OR^3,
4) $C(O)R^3$,
5) $C(O)N(R^3)_2$, or
6) $N(R^3)_2$;

Z is:

1) hydrogen,
2) R^6,
3) OR^3,
4) $N(R^3)_2$,
5) $C(O)OR^3$,
6) $C(O)N(R^3)_2$,
7) $C(O)R^3$,
8) halogen,
9) $N(R^3)(CR^{1c}{}_2)_nC(O)N(R^3)_2$,
10) $S(O)_mN(R^3)_2$,
11) $N(R^3)C(O)OR^3$,
12) $N(R^3)S(O)_mR^6$,
13) $N(R^3)C(O)R^3$,
14) $N(R^3)(CR^{1c}{}_2)_nR^3$, or
15) $S(O)_mR^6$;

n is independently 0 to 4;

or a pharmaceutically acceptable salt or stereoisomer thereof.

4. (Original) A compound selected from:

5-Chloro-3-[(methylamino)sulfonyl]-1*H*-indole-2-carboxamide;

3-(Aminosulfonyl)-5-chloro-1*H*-indole-2-carboxamide;

5-Bromo-3-({methyl[(5-oxo-4,5-dihydro-1H-1,2,4-triazol-3 yl)methyl] amino} sulfonyl)-1H-indole-2-carboxamide;

3-({[2-(Aminosulfonyl)ethyl]amino}sulfonyl)-5-iodo-1*H*-indole-2-carboxamide;
3-[(Dimethylamino)sulfonyl]-5-methoxy-1H-indole-2-carboxamide;

5-Chloro-3-{[(2-phenethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Chloro-3-[(benzylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-[(cyclohexylamino)sulfonyl]-1*H*-indole-2-carboxamide ;

5-Chloro-3-[(1-naphthylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-{[(3-phenylpropyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Chloro-3-[(ethylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-[(propylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-[(butylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-[(pentylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-{[ethyl(methyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Chloro-3-[(diethylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-[(*iso*-propylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-[(cyclobutylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-[(cyclopentylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-{[(4-chlorophenyl)amino}sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-{[(3-chlorophenyl)amino}sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-{[(2-chlorophenyl)amino}sulfonyl]-1*H*-indole-2-carboxamide;
5-Chloro-3-{[(4-chlorophenyl)methylamino}sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-{[(3-chlorophenyl)methylamino}sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-{[(2-chlorophenyl)methylamino}sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-[(*tert*-butylamino)sulfonyl]-1*H*-indole-2-carboxamide;

(±)-5-Chloro-3-[(pyrrolidin-3-ylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-[(piperidin-4-ylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-{[(1-methyl-1*H*-benzimidazol-2-yl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Chloro-3-[(benzamideamino)sulfonyl]-1*H*-indole-2-carboxamide;
5-Chloro-3-[(5-aminotetrazole)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-[(pyridin-4-ylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-[(pyridin-2-ylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Chloro-3-{[(2-methyoxyethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Chloro-3-[(dimethylamino)sulfonyl]-1*H*-indole-2-carboxamide;

3-({[2-(Aminosulfonyl)ethyl]amino}sulfonyl)-5-chloro-1*H*-indole-2-carboxamide;

5-Chloro-3-{[(2-hydroxyethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Chloro-3-{[(2-morpholin-4-ylethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Chloro-3-{[(2-methoxyethyl)(methyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-[({[2-(2-acetamide)amino]ethyl}amino)sulfonyl]-1*H*-indole-2-carboxamide;

N-{[2-(Aminocarbonyl)-5-bromo-1*H*-indol-3-yl]sulfonyl}-*N*-methyl-β-alaninamide;

5-Bromo-3-[(methylamino)sulfonyl]-1*H*-indole-2-carboxamide;

Ethyl *N*-{[2-(aminocarbonyl)-5-bromo-1*H*-indol-3-yl]sulfonyl} *N*-methyl-β-alaninate;

5-Bromo-3-{[cyclopropyl(methyl)amino]sulfonyl}-1H-indole-2-carboxamide;

(±)-5-Bromo-3-{[methyl(tetrahydrofuran-3-yl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-({methyl[2-(1*H*-1,2,4-triazol-1-yl)ethyl]amino}sulfonyl)-1*H*-indole-2-carboxamide;

5-Bromo-3-{[methyl(tetrahydro-2*H*-pyran-4-yl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

(±)-5-Bromo-3-{[(1,4-dioxan-2-ylmethyl)(methyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

3-({[4-(Aminosulfonyl)benzyl]amino}sulfonyl)-5-bromo-1*H*-indole-2-carboxamide;

5-Chloro-3-{[*iso*-propyl(2-methoxyethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

3-{[(2-Bromoethyl)(2-hydroxyethyl)amino]sulfonyl}-5-hydroxy-1*H*-indole-2-carboxamide;

3-{[(2-Bromoethyl)(2-hydroxyethyl)amino]sulfonyl}-5-methoxy-1*H*-indole-2-carboxamide;

5-Chloro-3-{[methoxy(methyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

(±)-5-Chloro-3-{[(2,3-dihydroxypropyl)(methyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Chloro-3-{[(2-hydroxyethyl)(methyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

N-{[2-(Aminocarbonyl)-5-chloro-1*H*-indol-3-yl]sulfonyl}-*N*-methylglycine;

N-{[2-(Aminocarbonyl)-5-chloro-1*H*-indol-3-yl]sulfonyl}-*N*-methylglycinamide;

5-Bromo-3-({[4-(methylsulfonyl)benzyl]amino}sulfonyl)-1*H*-indole-2-carboxamide;

3-[({2-[4-(Aminosulfonyl)phenyl]ethyl}amino)sulfonyl]-5-bromo-1*H*-indole-2-carboxamide;

3-{[(5-Amino-5-oxopentyl)amino]sulfonyl}-5-bromo-1*H*-indole-2-carboxamide;

3-({[2-(Aminosulfonyl)ethyl]amino}sulfonyl)-5-bromo-1*H*-indole-2-carboxamide;

tert-Butyl 2-({[2-(aminocarbonyl)-5-bromo-1*H*-indol-3-yl]sulfonyl}amino)-ethylcarbamate;

3-{[(2-Aminoethyl)amino]sulfonyl}-5-bromo-1*H*-indole-2-carboxamide;

5-Bromo-3-[({ethylsulfonylamino}ethylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Iodo-3-{[(2-{[(4-methoxyphenyl)sulfonyl]amino}ethyl)amino]sulfonyl}-1 *H*-indole-2-carboxamide;

5-Bromo-3-{[methoxy(methyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Fluoro-3-{[(2-{[(4-methoxyphenyl)sulfonyl]amino}ethyl)(methyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-{[(2-{[(4-nitrophenyl)sulfonyl]amino}ethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-({[2-({[(4-methoxyphenyl)amino]carbonyl}amino)ethyl]amino}sulfonyl)-1*H*-indole-2-carboxamide;

5-Bromo-3-[({3-[(4-chlorophenyl)thio]propyl}amino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Bromo-3-[({3-[(4-chlorophenyl)thio]propyl}amino)sulfonyl]-1 *H*-indole-2-carboxamide;

5-Bromo-3-[({3-[(4-chlorophenyl)sulfonyl]propyl}amino)sulfonyl]-1 *H*-indole-2-carboxamide;

5-Bromo-3-[({propylsulfonylamino}ethylamino)sulfonyl]-1*H*-indole-2-carboxamide hydrochloride;

5-Bromo-3-{[(2-{[(4-methoxyphenyl)sulfonyl]amino}ethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide ;

5-Bromo-3-[({2-[(phenylsulfonyl)amino]ethyl}amino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Bromo-3-[({2-[(methylsulfonyl)amino]ethyl}amino)sulfonyl]-1*H*-indole-2-carboxamide;

3-[({2-[(Benzylsulfonyl)amino]ethyl}amino)sulfonyl]-5-bromo-1*H*-indole-2-carboxamide;

5-Bromo-3-{[(2-{[(3-methoxyphenyl)sulfonyl]amino}ethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-{[(2-{[(2,5-dimethoxyphenyl)sulfonyl]amino}ethyl)amino]sulfonyl}-1*H* -indole-2-carboxamide;
5-Bromo-3-{[(2-{[(5-bromo-2-methoxyphenyl)sulfonyl]amino}ethyl)amino] sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-({[2-({[2-(trifluoromethoxy)phenyl]sulfonyl}amino)ethyl]amino} sulfonyl)-1 *H*-indole-2-carboxamide;

5-Bromo-3-{[(2-{[(2-methoxy-5-methylphenyl)sulfonyl]amino}ethyl)amino] sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-{[(2-{[(4-cyanophenyl)sulfonyl]amino}ethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-{[(2-{[(4-chlorophenyl)sulfonyl]amino}ethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-{[(2-{[(3,4-dimethoxyphenyl)sulfonyl]amino}ethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-[({3-[(phenylsulfonyl)amino]propyl}amino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Bromo-3-{[(3-{[(4-methoxyphenyl)sulfonyl]amino}propyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;
3-[({3-[(Benzylsulfonyl)amino]propyl}amino)sulfonyl]-5-bromo-1*H*-indole-2-carboxamide;

3-[({2-[(Aminocarbonyl)amino]ethyl}amino)sulfonyl]-5-bromo-1*H*-indole-2-carboxamide;

5-Bromo-3-{[(2-{[(4-bromophenyl)sulfonyl]amino}ethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-[({2-[(thien-3-ylsulfonyl)amino]ethyl}amino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Bromo-3-{[(2-{[(3-chlorobenzyl)sulfonyl]amino}ethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-{[(2-{[(2-phenylethyl)sulfonyl]amino}ethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-[({2-[(4-methoxybenzoyl)amino]ethyl}amino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Bromo-3-[({2-[(4-methoxybenzyl)amino]ethyl}amino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Bromo-3-[({2-[(4-methoxyphenyl)amino]ethyl}amino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Bromo-3-[({2-[(4-methoxyphenyl)(methylsulfonyl)amino]ethyl}amino)sulfonyl]-1*H*-indole-2-carboxamide;

3-[({2-[Acetyl(4-methoxyphenyl)amino]ethyl}amino)sulfonyl]-5-bromo-1*H*-indole-2-carboxamide;

5-Iodo-3-{[cyclopropyl(methyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Iodo-3-[(cyclopropylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Bromo-3-[(cyclopropylamino)sulfonyl]-1*H*-indole-2-carboxamide;

5-Iodo-3-{[methoxy(methyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

(±)-5-Chloro-3-{[(tetrahydro-2*H*-pyran-2-ylmethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

(±)-5-Bromo-3-{[(tetrahydro-2*H*-pyran-2-ylmethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

(±)-5-Iodo-3-{[(tetrahydro-2*H*-pyran-2-ylmethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

(±)-5-Chloro-3-{[methyl(tetrahydro-2*H*-pyran-2-ylmethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;
(±)-5-Bromo-3-{[methyl(tetrahydro-2*H*-pyran-2-ylmethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;
(±)-5-Iodo-3-{[methyl(tetrahydro-2*H*-pyran-2-ylmethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

5-Bromo-3-({[2-(tert-butylthio)ethyl]amino}sulfonyl)-1-*H*-indole-2-carboxamide;

5-chloro-3-{[methyl(tetrahydro-2H-pyran-4-yl)amino]sulfonyl}-1H-indole-2-carboxamide;

5-chloro-3-({[1-(2,3-dihydro-1,4-benzodioxin-2-yl)ethyl]amino}sulfonyl)-1H-indole-2-carboxamide;
5-chloro-3-[(tetrahydro-2H-pyran-4-ylamino)sulfonyl]-1H-indole-2-carboxamide;

5-chloro-3-{[(1,4-dioxan-2-ylmethyl)(methyl)amino]sulfonyl}-1H-indole-2-carboxamide;

5-chloro-3-({[(3-methyloxetan-3-yl)methyl]amino}sulfonyl)-1H-indole-2-carboxamide;

5-chloro-3-[(tetrahydrofuran-3-ylamino)sulfonyl]-1H-indole-2-carboxamide;

5-chloro-3-({[(1,1-dioxidotetrahydrothien-3-yl)methyl]amino}sulfonyl)-1H-indole-2-carboxamide;

5-chloro-3-({[2-(3-phenyl-1*H*-1,2,4-triazol-5-yl)ethyl]amino}sulfonyl)-1*H*-indole-2-carboxamide;

5-chloro-3-({[2-(2-methoxyphenyl)ethyl]amino}sulfonyl)-1H-indole-2-carboxamide;

5-chloro-3-({[3-(trifluoromethyl)benzyl]amino}sulfonyl)-1H-indole-2-carboxamide;

5-chloro-3-({[2-(2,3-dihydro-1*H*-indol-1-yl)ethyl]amino}sulfonyl)-1*H*-indole-2-carboxamide;

5-chloro-3-({methyl[(1-methylpiperidin-3-yl)methyl]amino}sulfonyl)-1*H*-indole-2-carboxamide;

5-chloro-3-{[(2,3-dihydro-1,4-benzodioxin-2-ylmethyl) amino]sulfonyl}-1H-indole-2-carboxamide;

5-bromo-3-{[(3-ethoxypropyl) amino]sulfonyl}-1H-indole-2-carboxamide;

3-[({[2-(aminocarbonyl)-5-bromo-1H-indol-3-yl]sulfonyl}amino) methyl]-1-benzylpyrrolidine;

5-bromo3-({[(1-benzylpyrrolidin-3-yl)methyl]amino}sulfonyl)-1*H*-indole-2-carboxamide;

5-bromo-3-{[(3-pyridin-3-ylpropyl)amino]sulfonyl}-1*H*-indole-2-carboxamide;

1-[2-({[2-(aminocarbonyl)-5-bromo-1H-indol-3-yl]sulfonyl}amino)ethyl]-4-phenylpiperidine;

5-bromo-3-{[(3-cyclohexylpropyl)amino]sulfonyl}-1H-indole-2-carboxamide;

5-bromo-3-{[(4,4-diphenylbutyl)amino]sulfonyl}-1H-indole-2-carboxamide;

5-bromo-3-{[(3-butoxypropyl)amino]sulfonyl}-1H-indole-2-carboxamide;

5-bromo-3-{[(6,7,8,9-tetrahydro-5H-benzo[a][7]annulen-7-ylmethyl)amino]sulfonyl}-1H-indole-2-carboxamide;

5-bromo-3-({[3-(3,5-dimethyl-1H-pyrazol-1-yl)propyl]amino}sulfonyl)-1H-indole-2-carboxamide;

5-bromo-3-({[3-(4-tert-butoxyphenyl)propyl]amino} sulfonyl)-1H-indole-2-carboxamide;

5-bromo-3-({[4-(4-tert-butoxyphenyl)butyl]amino}sulfonyl)-1H-indole-2-carboxamide;

5-bromo-3-{[(2-methoxy-1-methylethyl)amino]sulfonyl}-1H-indole-2-carboxamide;

5-bromo-3-{[(4-phenylbutyl)amino]sulfonyl}-1H-indole-2-carboxamide;

5-bromo-3-[({2-[(2,6-dichlorobenzyl)thio]ethyl}amino) sulfonyl]-1H-indole-2-carboxamide;

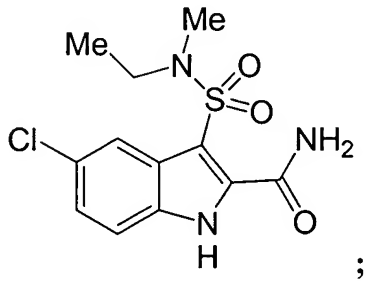
5-bromo-3-({[2-(tert-butylthio)ethyl]amino}sulfonyl)-1H-indole-2-carboxamide;

5-bromo-3-[({6-[(4-chlorobenzyl)amino]-6-oxohexyl}amino)sulfonyl]-1H-indole-2-carboxamide;

or a pharmaceutically acceptable salt or stereoisomer thereof.

5. (Original) The compound according to Claim 4, that is selected from:

5-Chloro-3-{[ethyl(methyl)amino]sulfonyl}-1*H*-indole-2-carboxamide



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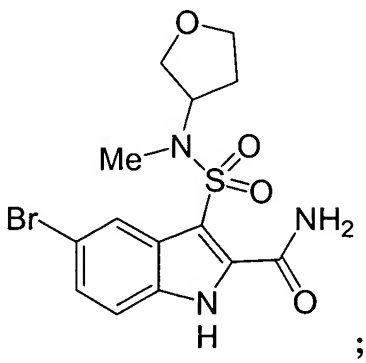
(±)-5-Bromo-3-{[methyl(tetrahydrofuran-3-yl)amino]sulfonyl}-1*H*-indole-2-carboxamide



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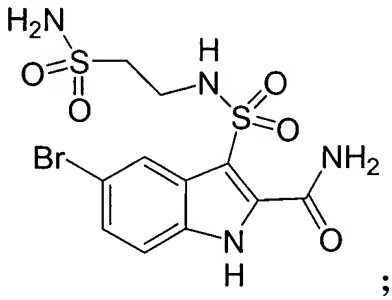
3-({[2-(Aminosulfonyl)ethyl]amino}sulfonyl)-5-bromo-1*H*-indole-2-carboxamide



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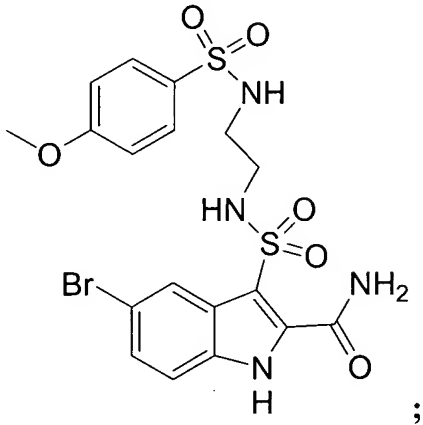
5-Bromo-3-{[(2-{[(4-methoxyphenyl)sulfonyl]amino}ethyl)amino]sulfonyl}-1*H*-indole-2-carboxamide



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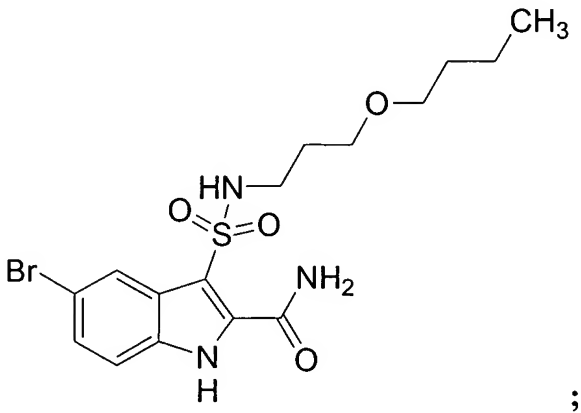
5-bromo-3-{[(3-butoxypropyl)amino]sulfonyl}-1H-indole-2-carboxamide



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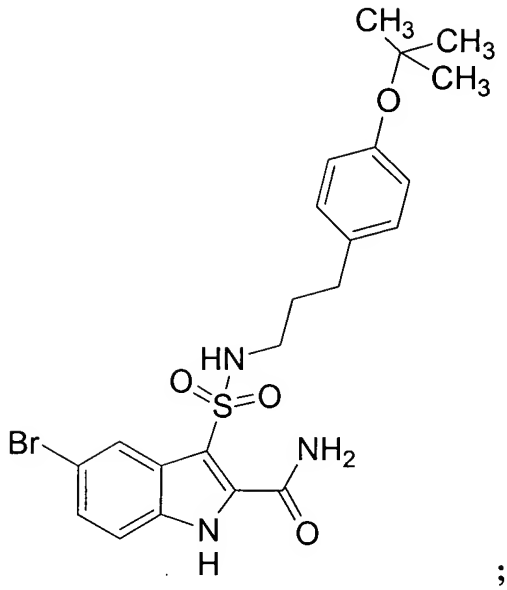
5-bromo-3-({[3-(4-tert-butoxyphenyl)propyl]amino}sulfonyl)-1H-indole-2-carboxamide



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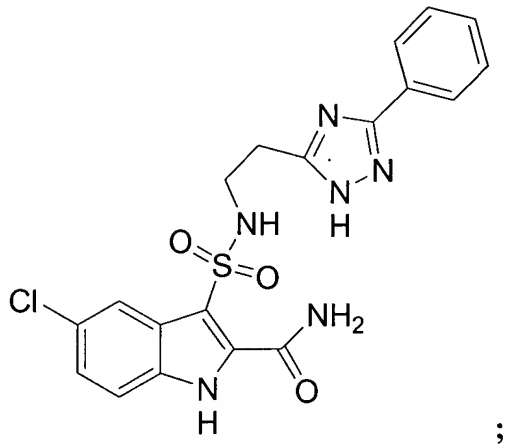
5-chloro-3-({[2-(3-phenyl-1*H*-1,2,4-triazol-5-yl)ethyl]amino}sulfonyl)-1*H*-indole-2-carboxamide



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or a pharmaceutically acceptable salt or stereoisomer thereof.

6. (Original) A pharmaceutical composition which is comprised of a compound in accordance with Claim 1 and a pharmaceutically acceptable carrier.

7. (Original) A method of modulating the catalytic activity of protein kinases in a mammal in need thereof comprising contacting the protein kinase with a compound of Claim 1.

8. (Original) The method of Claim 7 wherein the protein kinase is an RTK.

9. (Original) The method of Claim 8, wherein the RTK is selected from IR, IGF-1R and IRR.

10. (Original) A method of treating or preventing a PK-related disorder in a mammal in need thereof comprising administering to said mammal a therapeutically effective amount of a compound of Claim 1.

11. (Original) A method of Claim 10, wherein the PK-related disorder is an IGF-1R-related disorder selected from:

1) cancer,
2) diabetes,
3) an autoimmune disorder,
4) a hyperproliferation disorder,
5) aging,
6) acromegaly, and
7) Crohn's disease.

12. (Original) A method of treating cancer in a mammal in need of such treatment comprising administering to said mammal a therapeutically effective amount of a compound of Claim 1.

13. (Original) A method of treating retinal vascularization comprising administering to a mammal in need of such treatment a therapeutically effective amount of a compound of Claim 1.

14. (Original) A method of treating cancer which comprises administering a therapeutically effective amount of a compound of Claim 1 in combination with a second

compound selected from:

1) an estrogen receptor modulator,
2) an androgen receptor modulator,
3) retinoid receptor modulator,
4) a cytotoxic agent,
5) an antiproliferative agent,
6) a prenyl-protein transferase inhibitor,
7) an HMG-CoA reductase inhibitor,
8) an HIV protease inhibitor,
9) a reverse transcriptase inhibitor, and
10) an angiogenesis inhibitor.

15. (Original) The method of Claim 14, wherein the second compound is an estrogen receptor modulator selected from tamoxifen and raloxifene.

16. (Original) A method of treating cancer which comprises administering a therapeutically effective amount of a compound of Claim 1 in combination with radiation therapy.

17. (Original) The method of Claim 16 wherein radiation therapy is also administered.

18. (Original) A method of treating cancer which comprises administering a therapeutically effective amount of a compound of Claim 1 and paclitaxel or trastuzumab.

19. (Original) A method of treating or preventing cancer which comprises administering a therapeutically effective amount of a compound of Claim 1 and a GPIIb/IIIa antagonist.

20. (Canceled)

21. (Canceled)

22. (Original) A process for preparing an alkyl 5-iodo-1*H*-indole-2-carboxylate which comprises the steps of:

a) combining alkyl 1*H*-indole-2-carboxylate, iodine, sodium periodate and sulfuric acid in an alcohol, and heating to a temperature of about 50 °C to about 100 °C to obtain a product;
b) adding the product to a solution of organic solvent and aqueous solution to create a first biphasic mixture;
c) removing, drying, filtering and concentrating the organic layer;
d) dissolving the organic layer in an alcohol;
e) adding zinc and aqueous acid to produce a mixture;
f) combining the mixture with water to create a second biphasic mixture; and
g) extracting, drying and filtering the organic layer of the second biphasic mixture to obtain the alkyl 5-iodo-1*H*-indole-2-carboxylate.

23. (Original) The process of Claim 22 wherein the alkyl 5-iodo-1*H*-indole-2-carboxylate is ethyl 5-iodo-1*H*-indole-2-carboxylate.